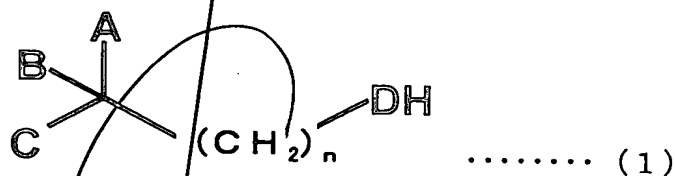


CLAIMS

1. A process for producing optically active 4-amino-2-methylbutane-1-ol which comprises: treating racemic 4-amino-2-methylbutane-1-ol with an optically active organic acid to obtain a diastereomeric salt, crystallizing out the resulting diastereomeric salt, and subjecting the salt to solid-liquid separation.
2. The process for producing optically active 4-amino-2-methylbutane-1-ol according to claim 1 wherein the optically active organic acid is an optically active carboxylic acid, optically active sulfonic acid or optically active phosphonic acid represented by the formula (1)



- wherein D denotes COO^- , SO_3^- or PO_3H^- ; A, B and C each denote hydrogen, a substituted or unsubstituted, straight or branched chain alkyl group having 1-10 carbon atoms, halogen atom, alkoxy group, hydroxyl group, nitro group, carboxyl group, acyloxy group, or substituted or unsubstituted amino group, phenyl group or naphthyl group; the substituent in the alkyl group, amino group, phenyl group or naphthyl group is a straight or branched chain alkyl group having 1-10 carbon atoms, halogen atom, alkoxy group, hydroxyl

group, nitro group, benzoyl group, carboxyl group, acyl group, methylthio group or sulfonic acid group; provided that A, B, C and $(CH_2)_n-DH$ are not the same with each other at the same time; and n is 1 or 0.

- 5 3. The process according to claim 1 wherein the optically active organic acid is an optically active 2-aryl-2-substituted acetic acid represented by the following formula (2)

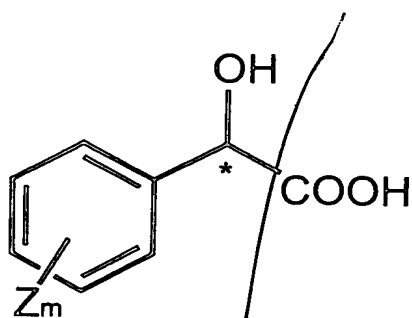


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wherein Y denotes a straight or branched chain alkyl group having 1-10 carbon atoms, halogen atom, alkoxy group, acyloxy group or hydroxyl group; Ar denotes a substituted or unsubstituted phenyl group or naphthyl group; the substituent is a straight or branched chain alkyl group having 1-10 carbon atoms, halogen atom, alkoxy group, hydroxyl group, nitro group, benzoyl group, carboxyl group, methylthio group or sulfonic acid group; and * denotes asymmetric carbon.

- 20 4. The process according to claim 1 wherein the optically active organic acid is an optically active mandelic acid derivative represented by the following formula (3)

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..... (3)

wherein Z denotes hydrogen or a straight or branched chain alkyl group having 1-10 carbon atoms, halogen atom, alkoxy group, hydroxyl group, nitro group, methylthio group or benzoyl group; * denotes asymmetric carbon; m is an integer of from 1 to 5 and; when $m \geq 2$, Z may be same as or different from each other.

5. The process according to any one of claims 1-4 wherein the racemic 4-amino-2-methylbutane-1-ol is mixed with the optically active organic acid by use of a solvent, and mother liquor is removed at the time of the solid-liquid separation.

6. The process according to claim 5 wherein the solvent is at least one component selected from the group consisting of water, methanol, ethanol, isopropanol, n-propanol, isopropyl ether, acetone and acetonitrile.

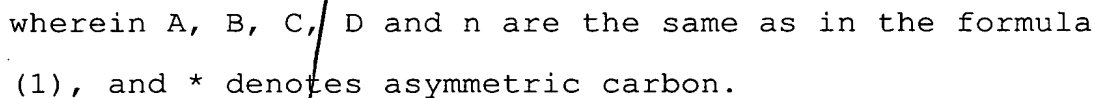
7. The process according to any one of claims 1-6 wherein the diastereomeric salt obtained is recrystallized by using a solvent to obtain a diastereomeric salt of higher optical purity.

8. The process according to claim 7 wherein the

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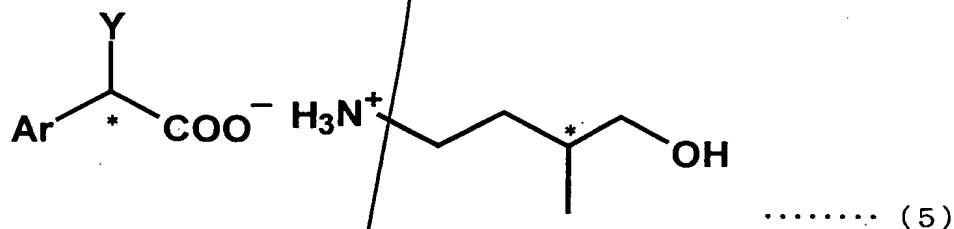
9. The process according to any one of claims 1-8 wherein optically active 4-amino-2-methylbutane-1-ol is obtained by neutralizing the diastereomeric salt obtained or passing the salt through ion exchange resin.

11. The salt of optically active 4-amino-2-methylbutane-1-ol with an optically active 4-amino-2-methylbutane-1-ol according to claim 10 wherein the optically active organic acid is an optically active carboxylic acid, optically active sulfonic acid or optically active phosphonic acid and the structure of the salt is represented by the formula (4)



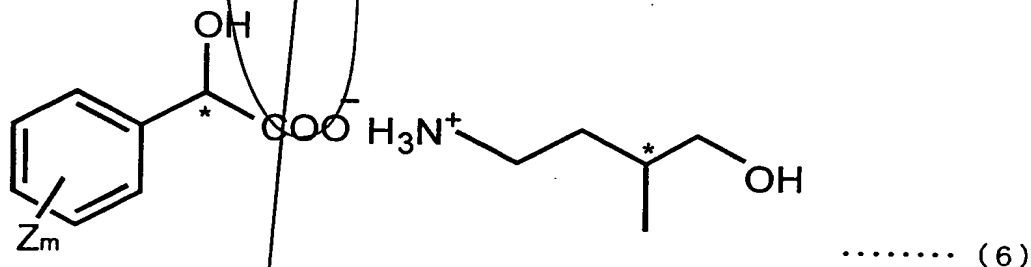
12. The salt of optically active 4-amino-2-

methylobutane-1-ol with an optically active organic acid according to claim 10 wherein the optically active organic acid is an optically active 2-aryl-2-substituted acetic acid and the structure of the salt is represented by the formula (5)



wherein Ar and Y are the same as in the formula (2), and * denotes asymmetric carbon.

13. The salt of optically active 4-amino-2-methylbutane-1-ol with an optically active organic acid according to claim 10 wherein the optically active organic acid is an optically active mandelic acid derivative and the structure of the salt is represented by the formula (6)



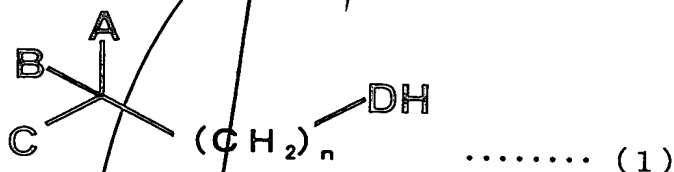
wherein Z and m are the same as in the formula (3).

14. A process for producing a salt of optically

active 4-amino-2-methylbutane-1-ol with an optically active organic acid which comprises; treating racemic 4-amino-2-methylbutane-1-ol with an optically active organic acid to obtain a diastereomeric salt,

- 5 crystallizing out the resulting diastereomeric salt, and subjecting the salt to solid-liquid separation.

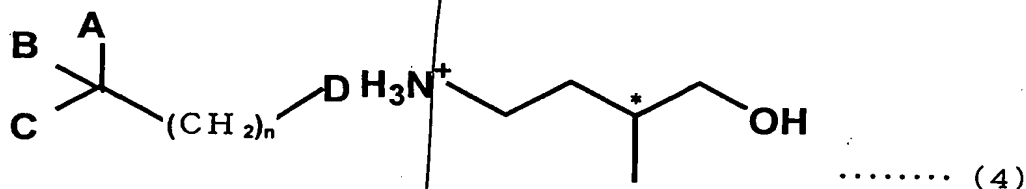
15. The process according to claim 14 wherein the optically active organic acid is an optically active carboxylic acid, optically active sulfonic acid or
10 optically active phosphonic acid represented by the formula (1) and the structure of the salt obtained is represented by the formula (4)



wherein D denotes COO^- , SO_3^- or PO_3H^- ; A, B and C each
15 denote hydrogen, a substituted or unsubstituted, straight or branched chain alkyl group having 1-10 carbon atoms, halogen atom, alkoxy group, hydroxyl group, nitro group, carboxyl group, acyloxy group, or substituted or unsubstituted amino group, phenyl group
20 or naphthyl group; the substituent in said alkyl group, amino group, phenyl group or naphthyl group is a straight or branched chain alkyl group having 1-10 carbon atoms, halogen atom, alkoxy group, hydroxyl group, nitro group, benzoyl group, carboxyl group, acyl

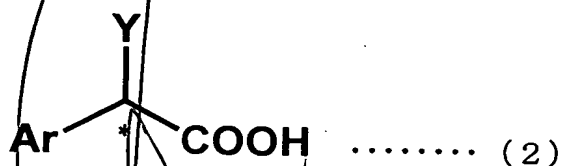
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group, methylthio group or sulfonic acid group;
provided that A, B, C and $(CH_2)_n-DH$ are not the same
with each other at the same time; and n is 1 or 0,



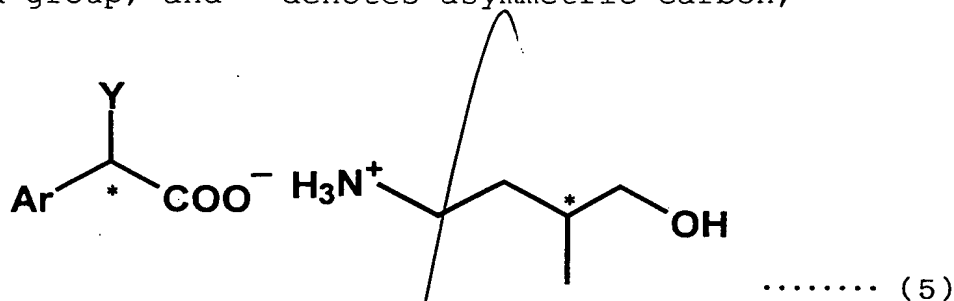
5 wherein A, B, C, D and n are the same as in the formula
(1), and * denotes asymmetric carbon.

16. The process according to claim 14 wherein the
optically active organic acid is an optically active 2-
aryl-2-substituted acetic acid represented by the
10 following formula (2), and the structure of the salt
obtained is represented by the formula (5)



wherein Y denotes a straight or branched chain alkyl
group having 1 to 10 carbon atoms, halogen atom, alkoxy
15 group, acyloxy group or hydroxyl group; Ar denotes a
substituted or unsubstituted phenyl group or naphthyl
group; the substituent is a straight or branched chain
alkyl group having 1-10 carbon atoms, halogen atom,
alkoxy group, hydroxyl group, nitro group, benzoyl
20 group, carboxyl group, methylthio group or sulfonic

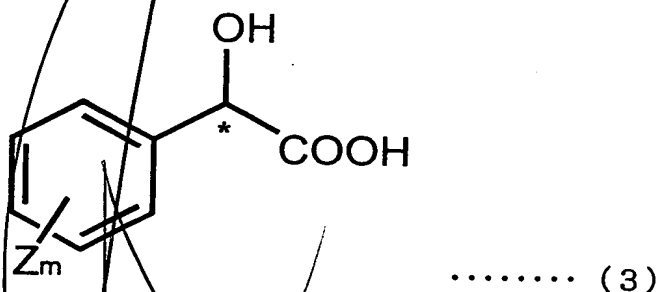
acid group; and * denotes asymmetric carbon,



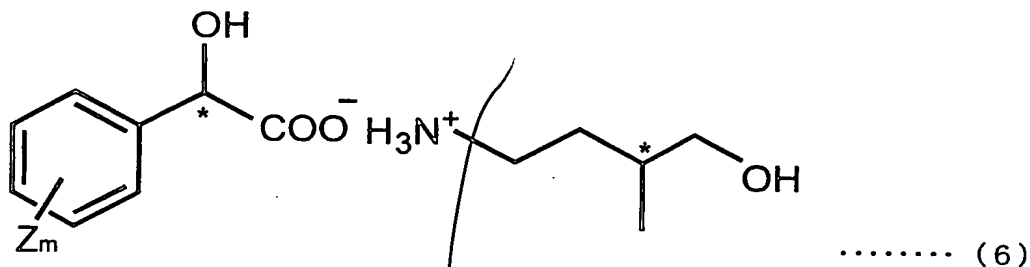
wherein Ar and Y are the same as in the formula (2),
and * denotes asymmetric carbon.

- 5 17. The process according to claim 14 wherein the
optically active organic acid is an optically active
mandelic acid derivative represented by the following
formula (3) and the structure of the salt obtained is
represented by the formula (6),

10



- wherein Z is hydrogen or a straight or branched chain
alkyl group having 1-10 carbon atoms, halogen atom,
alkoxy group, hydroxyl group, nitro group, methylthio
15 group or benzoyl group; * denotes asymmetric carbon; m
is an integer of from 1 to 5 and; when $m \geq 2$, Z may be
same as or different from each other,



wherein Z and m are the same as in the formula (3).

18. The process according to any one of claims 14-17 wherein the racemic 4-amino-2-methylbutane-1-ol is mixed with the optically active organic acid by use of a solvent, and mother liquor is removed at the time of the solid-liquid separation.

19. The process according to claim 18 wherein the solvent is at least one component selected from the group consisting of water, methanol, ethanol, isopropanol, n-propanol, isopropyl ether, acetone and acetonitrile.

20. The process according to any one of claims 14-19 wherein the diastereomeric salt obtained is recrystallized by using a solvent to obtain a diastereomeric salt of higher optical purity.

21. The process according to claim 20 wherein the solvent used for the recrystallization is at least one component selected from the group consisting of water, methanol, ethanol, isopropanol, n-propanol, isopropyl ether, acetone and acetonitrile.

22. A process for producing optically active 4-amino-2-methylbutane-1-ol by using the salt according

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to any one of claims 10-13.

23. A process for producing optically active 4-amino-2-methylbutane-1-ol by using the salt obtained by the process according to any one of claims 14-21.

5 24. A process for producing optically active 4-amino-2-methylbutane-1-ol which comprises; bringing a diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution into contact with a solvent and an
10 alkali to decompose the salt, subjecting the resulting reaction mixture to solid-liquid separation to obtain a filtrate, and obtaining optically active 4-amino-2-methylbutane-1-ol from the filtrate.

25. The process for producing optically active 4-amino-2-methylbutane-1-ol according to claim 24 wherein
15 a filtration residue containing an alkali salt of the optically active reagent for optical resolution is obtained by the solid-liquid separation, the filtration residue is brought into contact with a solvent and an
20 acid, and the optically active reagent for optical resolution thus crystallized out is subjected to solid-liquid separation and recovered.

26. The process for producing optically active 4-amino-2-methylbutane-1-ol according to claim 24 wherein
25 the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is brought into contact with a solvent and an alkali to decompose the salt, the

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solvent is replaced with an alcohol in which a solubility of an alkali salt of the optically active reagent for optical resolution is low, and an alkali salt of the optically active reagent for optical resolution and the optically active 4-amino-2-methylbutane-1-ol solution are subjected to solid-liquid separation to recover the alkali salt of the optically active reagent for optical resolution.

27. The process for producing optically active 4-amino-2-methylbutane-1-ol according to claim 24 wherein the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is brought into contact with an alcohol and an alkali metal alcoholate to decompose the salt, the alcohol is replaced with an alcohol in which a solubility of an alkali metal salt of the optically active reagent for optical resolution is low, and an alkali metal salt of the optically active reagent for optical resolution and an optically active 4-amino-2-methylbutane-1-ol solution are subjected to solid-liquid separation to recover the alkali metal salt of the optically active reagent for optical resolution.

28. The process for producing optically active 4-amino-2-methylbutane-1-ol according to claim 24 wherein the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is brought into contact with water and/or an alcohol and an alkali metal hydroxide to

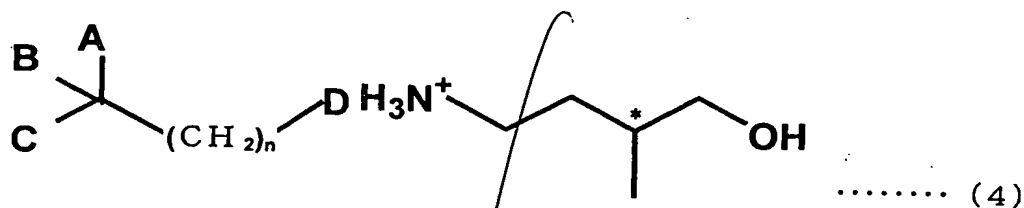
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decompose the salt, the water and/or the alcohol are replaced with an alcohol in which a solubility of an alkali metal salt of the optically active reagent for optical resolution is low, and an alkali metal salt of the optically active reagent for optical resolution and an optically active 4-amino-2-methylbutane-1-ol solution are subjected to solid-liquid separation to recover the alkali metal salt of the optically active reagent for optical resolution.

29. The process for producing optically active 4-amino-2-methylbutane-1-ol according to any one of claims 24-28 wherein the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is a salt of optically active 4-amino-2-methylbutane-1-ol with an optically active organic acid.

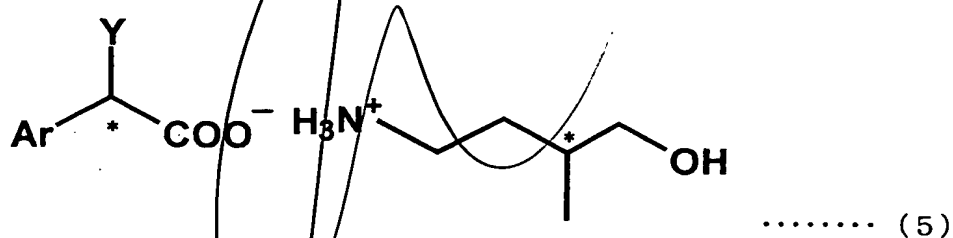
30. The process for producing optically active 4-amino-2-methylbutane-1-ol according to any one of claims 24-28 wherein the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is a salt of optically active 4-amino-2-methylbutane-1-ol with an optically active carboxylic acid, optically active sulfonic acid or optically active phosphonic acid represented by the following formula (4)

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wherein A, B, C, D and n are the same as in the formula (1), and * denotes asymmetric carbon.

31. The process for producing optically active 4-amino-2-methylbutane-1-ol according to any one of claims 24-28 wherein the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is a salt of optically active 4-amino-2-methylbutane-1-ol with an optically active 2-aryl-2-substituted acetic acid represented by the following formula (5)

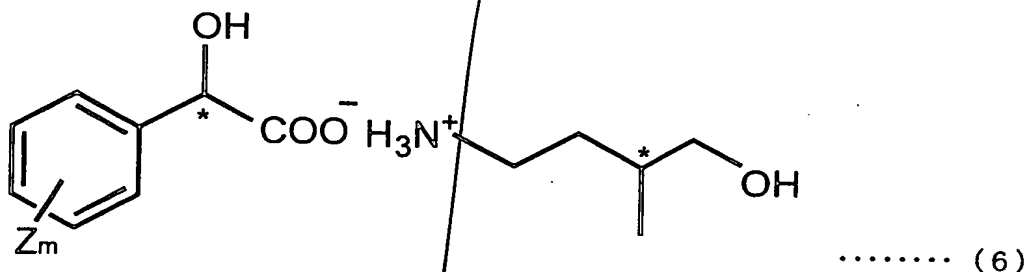


wherein Ar and Y are the same as in the formula (2), and * denotes asymmetric carbon.

32. The process for producing optically active 4-amino-2-methylbutane-1-ol according to any one of claims 24-28 wherein the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is a

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salt of optically active 4-amino-2-methylbutane-1-ol with an optically active mandelic acid derivative represented by the formula (6)



5

wherein Z and m are the same as in the formula (3).

33. A process for recovering an optically active reagent for optical resolution used in producing optically active 4-amino-2-methylbutane-1-ol which comprises: bringing a diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active optically resolving agent into contact with a solvent and an alkali to decompose the salt, subjecting the resulting reaction mixture to solid-liquid separation to obtain a filtration residue containing an alkali salt of the optically active reagent for optical resolution, bringing the filtration residue into contact with a solvent and an acid to crystallize out an optically active reagent for optical resolution, and subjecting the optically active reagent for optical resolution thus crystallized out to solid-liquid separation to recover it.

20

34. The recovering process according to claim 33

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wherein the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is brought into contact with a solvent and an alkali to decompose the salt, the solvent is replaced with an alcohol in which a solubility of an alkali salt of the optically active reagent for optical resolution is low, and an alkali salt of the optically active reagent for optical resolution and an optically active 4-amino-2-methylbutane-1-ol solution are subjected to solid-liquid separation to recover the alkali salt of the optically active reagent for optical resolution.

35. The recovering process according to claim 33 wherein the diastereomeric salt of optically active 4-amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is brought into contact with an alcohol and an alkali metal alcoholate to decompose the salt, the alcohol is replaced with an alcohol in which a solubility of an alkali metal salt of the optically active reagent for optical resolution is low, and an alkali metal salt of the optically active reagent for optical resolution and an optically active 4-amino-2-methylbutane-1-ol solution are subjected to solid-liquid separation to recover the alkali metal salt of the optically active reagent for optical resolution.

36. The recovering process according to claim 33 wherein the diastereomeric salt of optically active 4-

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amino-2-methylbutane-1-ol and an optically active reagent for optical resolution is brought into contact with water and/or alcohol and an alkali metal hydroxide to decompose the salt, the water and/or alcohol are

5 replaced with an alcohol in which a solubility of an alkali metal salt of the optically active reagent for optical resolution is low, and an alkali metal salt of the optically active reagent for optical resolution and an optically active 4-amino-2-methylbutane-1-ol
10 solution are subjected to solid-liquid separation to recover the alkali metal salt of the optically active reagent for optical resolution.

37. The recovering process according to any one of claims 33-36 wherein the optically active reagent
15 for optical resolution is an optically active organic acid.

38. The recovering process according to any one of claims 33-36 wherein the optically active reagent for optical resolution is an optically active
20 carboxylic acid, optically active sulfonic acid or optically active phosphonic acid represented by the above-mentioned formula (1).

39. The recovering process according to any one of claims 33-36 wherein the optically active reagent
25 for optical resolution is an optically active 2-aryl-2-substituted acetic acid represented by the above-mentioned formula (2).

40. The recovering process according to any one

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of claims 33-36 wherein the optically active reagent for optical resolution is an optically active mandelic acid derivative represented by the above-mentioned formula (3).

- 5 41. A process for producing optically active 4-amino-2-methylbutane-1-ol by reusing the optically active reagent for optical resolution recovered by the recovering process according to any one of claims 33-40.
- 10 42. A process for using the optically active 4-amino-2-methylbutane-1-ol obtained by the process according to any one of claims 1-9, 24-32 or 41 as an intermediate in synthesizing optically active medicines or pesticides.
- 15 43. A process for producing optically active medicines or pesticides by using the optically active 4-amino-2-methylbutane-1-ol obtained by the process according to any one of claims 1-9, 24-32 or 41.

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